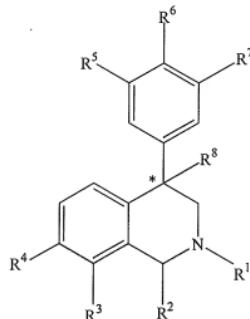


This listing of claims will replace all prior versions, and listings, of claims in the application:

Amendments to the Claims:

- (Previously presented) A method of treating urinary incontinence comprising administration of an effective amount of a compound of formula IA-IF having the following structure:



IA-IF

wherein:

the carbon atom designated \* is in the R or S configuration;

R<sup>1</sup> is C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl or C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl, each of which is optionally substituted with 1 to 3 substituents independently selected at each occurrence thereof from C<sub>1</sub>-C<sub>3</sub> alkyl, halogen, aryl, -CN, -OR<sup>9</sup> and -NR<sup>9</sup>R<sup>10</sup>;

R<sup>2</sup> is H, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl or C<sub>1</sub>-C<sub>6</sub> haloalkyl;

R<sup>3</sup> is H, halogen, -OR<sup>11</sup>, -S(O)R<sup>12</sup>, -S(O)<sub>n</sub> NR<sup>11</sup>R<sup>12</sup>, -CN, -C(O)R<sup>12</sup>, -C(O)NR<sup>11</sup>R<sup>12</sup>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl, -O(phenyl) or -O(benzyl), wherein each of -O(phenyl) and -O(benzyl) is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, or C<sub>1</sub>-C<sub>4</sub> alkoxy, or wherein R<sup>3</sup> is a C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl or C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl group, then said group is optionally substituted with from 1 to 3 substituents selected independently at each occurrence thereof from C<sub>1</sub>-C<sub>3</sub> alkyl, halogen, aryl, -CN, -OR<sup>9</sup> and -NR<sup>9</sup>R<sup>10</sup>,

provided that for compounds of formula IA, R<sup>3</sup> is C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl or C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl, each of which is optionally substituted with from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, or C<sub>1</sub>-C<sub>4</sub> alkoxy.

to 3 substituents selected independently at each occurrence thereof from C<sub>1</sub>-C<sub>3</sub> alkyl, halogen, aryl, -CN, -OR<sup>9</sup> and -NR<sup>9</sup>R<sup>10</sup>;

provided that for compounds of formula IB, R<sup>3</sup> is -O(phenyl), -O(benzyl), -OC(O)R<sup>13</sup> or -S(O)<sub>n</sub>R<sup>12</sup>, each of -O(phenyl) and -O(benzyl) is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, or C<sub>1</sub>-C<sub>4</sub> alkoxy;

R<sup>4</sup> is H, halogen, -OR<sup>11</sup>, -S(O)<sub>n</sub>R<sup>12</sup>, -S(O)NR<sup>11</sup>R<sup>12</sup>, -CN, -C(O)R<sup>12</sup>, -C(O)NR<sup>11</sup>R<sup>12</sup>, -NR<sup>11</sup>R<sup>12</sup>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl, O(phenyl) or -O(benzyl), wherein each of -O(phenyl) and -O(benzyl) is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl or C<sub>1</sub>-C<sub>4</sub> alkoxy and wherein R<sup>4</sup> is a C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl or C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl group, then said group is optionally substituted with from 1 to 3 substituents selected independently at each occurrence thereof from C<sub>1</sub>-C<sub>3</sub> alkyl, halogen, aryl, -CN, -OR<sup>9</sup> and -NR<sup>9</sup>R<sup>10</sup>,

provided that for compounds of formula IC, R<sub>4</sub> is C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, or C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl, each of which is optionally substituted with from 1 to 3 substituents selected independently at each occurrence thereof from C<sub>1</sub>-C<sub>3</sub> alkyl, halogen, aryl, -CN, -OR<sup>9</sup> and -NR<sup>9</sup>R<sup>10</sup>, or R<sup>5</sup> and R<sup>6</sup> or R<sup>6</sup> and R<sup>7</sup> may be -O-C(R<sup>12</sup>)<sub>2</sub>-O-;

provided that for compounds of formula ID, R<sup>4</sup> is -O(phenyl), -O(benzyl), -OC(O)R<sup>13</sup>, -NR<sup>11</sup>R<sup>12</sup> or -S(O)<sub>n</sub>R<sup>12</sup>, each of -O(phenyl) and -O(benzyl) is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, or C<sub>1</sub>-C<sub>4</sub> alkoxy;

R<sup>5</sup>, R<sup>6</sup> and R<sup>7</sup> in compounds of each of the formulae IA, IB, IC, ID, IE and IF are each independently H, halogen, -OR<sup>11</sup>, -S(O)<sub>n</sub>R<sup>12</sup>, -CN, -C(O)R<sup>12</sup>, -NR<sup>11</sup>R<sup>12</sup>, -C(O)NR<sup>11</sup>R<sup>12</sup>, -NR<sup>11</sup>C(O)R<sup>12</sup>, -NR<sup>11</sup>C(O)<sub>2</sub>R<sup>12</sup>, -NR<sup>11</sup>C(O)NR<sup>12</sup>R<sup>13</sup>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl or C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl, wherein each of R<sup>5</sup>, R<sup>6</sup> and R<sup>7</sup> is a C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl or C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl group, then said group is optionally substituted with from 1 to 3 substituents selected independently at each occurrence thereof from C<sub>1</sub>-C<sub>3</sub> alkyl, halogen, aryl, -CN, -OR<sup>9</sup> and -NR<sup>9</sup>R<sup>10</sup>, or R<sup>5</sup> and R<sup>6</sup> or R<sup>6</sup> and R<sup>7</sup> may be -O-C(R<sup>12</sup>)<sub>2</sub>-O-;

provided that for compounds of formula IE at least one of R<sup>5</sup> or R<sup>7</sup> is fluoro, chloro, or methyl;

or R<sup>7</sup> and R<sup>8</sup> are each independently -O-C(R<sup>12</sup>)<sub>2</sub>-0- in compounds of the formulae IE, but only where R<sup>2</sup> is fluoro, chloro or methyl;

or R<sup>7</sup> and R<sup>8</sup> can independently also be -O-C(R<sup>12</sup>)<sub>2</sub>-0- in compounds of the formulae IE, but only where R<sup>7</sup> is fluoro, chloro or methyl;

R<sup>8</sup> is H, halogen, or OR<sup>11</sup>, provided that for compounds of formula IF, R<sup>8</sup> is halogen;

R<sup>9</sup> and R<sup>10</sup> are each independently H, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>1</sub>-C<sub>4</sub> alkoxyalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl, -C(O)R<sup>13</sup>, phenyl or benzyl, where phenyl or benzyl is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, or C<sub>1</sub>-C<sub>4</sub> alkoxy;

or R<sup>9</sup> and R<sup>10</sup> are taken together with the nitrogen to which they are attached to form piperidine, pyrrolidine, piperazine, N-methylpiperazine, morpholine, or thiomorpholine; R<sup>11</sup> is H, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>1</sub>-C<sub>4</sub> alkoxyalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl, -C(O)R<sup>13</sup>, phenyl or benzyl, where R<sup>11</sup> is a C<sub>1</sub>-C<sub>4</sub> alkyl, phenyl or benzyl group, then said group is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, or C<sub>1</sub>-C<sub>4</sub> alkoxy;

R<sup>12</sup> is H, amino, C<sub>1</sub>-C<sub>4</sub> alkyl, (C<sub>1</sub>-C<sub>4</sub> alkyl)amino, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>1</sub>-C<sub>4</sub> alkoxyalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl, phenyl or benzyl, where phenyl or benzyl is optionally substituted from 1 to 3 times with a substituent selected independently from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl and C<sub>1</sub>-C<sub>6</sub> alkoxy;

or R<sup>11</sup> and R<sup>12</sup> are taken together with the nitrogen to which they are attached to form piperidine, pyrrolidine, piperazine, N-methylpiperazine, morpholine, or thiomorpholine; provided that only one of R<sup>9</sup> and R<sup>10</sup> or R<sup>9</sup> and R<sup>10</sup> are taken together with the nitrogen to which they are attached to form piperidine, pyrrolidine, piperazine, N-methylpiperazine, morpholine, or thiomorpholine;

R<sup>13</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl or phenyl;

n is 0, 1, or 2, and;

aryl is phenyl which is optionally substituted 1-3 times with halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl and C<sub>1</sub>-C<sub>4</sub> alkoxy,

or an oxide thereof, a pharmaceutically acceptable salt thereof, a solvate thereof, or prodrug thereof.

2. (Original) A method of claim 1, wherein R<sup>1</sup> is C<sub>1</sub>-C<sub>3</sub> alkyl.

3. (Original) A method of claim 2, wherein R<sup>1</sup> is CH<sub>3</sub>.
4. (Original) A method of claim 1, wherein R<sup>2</sup> is H, C<sub>1</sub>-C<sub>4</sub> alkyl or C<sub>1</sub>-C<sub>6</sub> haloalkyl.
5. (Original) A method of claim 4, wherein R<sup>2</sup> is H or CH<sub>3</sub>.
6. (Original) A method of claim 1, wherein R<sup>3</sup> is H or R<sup>3</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl or C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl, each of which is optionally substituted with from 1 to 3 substituents selected independently at each occurrence thereof from C<sub>1</sub>-C<sub>3</sub> alkyl, halogen, aryl, -CN, -OR<sup>9</sup> and NR<sup>9</sup>R<sup>10</sup>, or R<sup>3</sup> is -O(phenyl) or -O(benzyl) optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, or C<sub>1</sub>-C<sub>4</sub> alkoxy.
7. (Original) A method of claim 6, wherein R<sup>3</sup> is methyl, ethyl, propyl, or isopropyl.
8. (Original) A method of claim 6, wherein R<sup>3</sup> is -O(phenyl) or -O-CH<sub>2</sub>-(phenyl), each of which is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, or C<sub>1</sub>-C<sub>4</sub> alkoxy.
9. (Original) A method of claim 6, wherein R<sup>3</sup> is H.
10. (Original) A method of claim 1, wherein R<sup>4</sup> is H, or R<sup>4</sup> is -NR<sup>11</sup>R<sup>12</sup> or R<sup>4</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl or C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl, each of which is optionally substituted, or wherein R<sup>4</sup> is -O(phenyl) or -O(benzyl), each of which is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, or C<sub>1</sub>-C<sub>4</sub> alkoxy.
11. (Original) A method of claim 10, wherein R<sup>4</sup> is methyl, ethyl, propyl, or isopropyl.
12. (Original) A method of claim 10, wherein R<sup>4</sup> is -O(phenyl) or -O(CH<sub>2</sub>)phenyl, each of which is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, or C<sub>1</sub>-C<sub>4</sub> alkoxy.

13. (Original) A method of claim 10, wherein R<sup>4</sup> is H.
14. (Original) A method of claim 1, wherein R<sup>3</sup> and R<sup>4</sup> are each H or wherein R<sup>3</sup> and R<sup>4</sup> are each halogen.
15. (Original) A method of claim 1, wherein one of R<sup>3</sup> and R<sup>4</sup> is H and the other is CH<sub>3</sub>.
16. (Original) A method of claim 1, wherein R<sup>5</sup>, R<sup>6</sup> and R<sup>7</sup> are each H, halogen, -OR<sup>11</sup>, -NR<sup>11</sup>R<sup>12</sup>, C<sub>1</sub>-C<sub>6</sub> alkyl and substituted C<sub>1</sub>-C<sub>6</sub> alkyl.
17. (Original) A method of claim 16, wherein R<sup>5</sup>, R<sup>6</sup> and R<sup>7</sup> are each H.
18. (Original) A method of claim 16, wherein one of R<sup>5</sup> or R<sup>7</sup> is F, Cl or Me and the other of R<sup>5</sup> or R<sup>7</sup> and R<sup>8</sup> are H, halogen, -OR<sup>11</sup>, -NR<sup>11</sup>R<sup>12</sup>, or optionally substituted C<sub>1</sub>-C<sub>6</sub> alkyl.
19. (Original) A method of claim 18, wherein R<sup>5</sup> is F, Cl or Me; and R<sup>7</sup> is H.
20. (Original) The method of claim 18, wherein R<sup>5</sup> is F, Cl or Me; and R<sup>8</sup> is H.
21. (Original) A method of claim 1, wherein R<sup>8</sup> is halogen.
22. (Original) A method of claim 21, wherein R<sup>8</sup> is fluoro.
23. (Original) A method of claim 1, wherein:  
R<sup>1</sup> is C<sub>1</sub>-C<sub>3</sub> alkyl;  
R<sup>2</sup> is H, C<sub>1</sub>-C<sub>4</sub> alkyl or C<sub>1</sub>-C<sub>8</sub> haloalkyl;  
R<sup>3</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl or C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl, each of which is optionally substituted, or R<sup>3</sup> is -O(phenyl) or -O(benzyl), each of which is optionally substituted, or R<sup>3</sup> is H; R<sup>4</sup> is H, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl or C<sub>4</sub>-C<sub>7</sub> cycloalkylalkyl, each of which is optionally substituted with from 1 to 3 substituents selected independently at each occurrence thereof from C<sub>1</sub>-C<sub>3</sub> alkyl, halogen, aryl, -CN, -OR<sup>9</sup> and -NR<sup>9</sup>R<sup>10</sup>, or R<sup>4</sup> is -NR<sup>11</sup>R<sup>12</sup>, -O(phenyl) or -O(benzyl), wherein said -O(phenyl) or -O(benzyl), is optionally

substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, or C<sub>1</sub>-C<sub>4</sub> alkoxy;

or R<sup>3</sup> and R<sup>4</sup> are each halogen;

R<sup>5</sup>, R<sup>6</sup> and R<sup>7</sup> are each H, halogen, -OR<sup>11</sup>, -NR<sup>11</sup>R<sup>12</sup>, optionally substituted C<sub>1</sub>-C<sub>6</sub> alkyl, or one of R<sup>5</sup> and R<sup>7</sup> is Cl, F or Me and the other of R<sup>5</sup> and R<sup>7</sup> and R<sup>6</sup> is H, halogen, -OR<sup>11</sup>, -NR<sup>11</sup>R<sup>12</sup>, C<sub>1</sub>-C<sub>6</sub> alkyl or substituted C<sub>1</sub>-C<sub>6</sub> alkyl.

24. (Original) A method of claim 23, wherein:

R<sup>1</sup> is CH<sub>3</sub>;

R<sup>2</sup> is H or CH<sub>3</sub>;

R<sup>3</sup> is H, F, methyl, ethyl, propyl, isopropyl, -O(phenyl) or -0-CH<sub>2</sub>-(phenyl), wherein said -O(phenyl) or -0-CH<sub>2</sub>-(phenyl) is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, or C<sub>1</sub>-C<sub>4</sub> alkoxy;

R<sup>4</sup> is H, F methyl, ethyl, propyl, isopropyl, -O(phenyl) or -0-CH<sub>2</sub>-(phenyl), wherein said -O(phenyl) or -0-CH<sub>2</sub>-(phenyl) is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, or C<sub>1</sub>-C<sub>4</sub> alkoxy;

R<sup>5</sup>, R<sup>6</sup> and R<sup>7</sup> are each H or R<sup>5</sup> is F, Cl or Me, or one of R<sup>6</sup> or R<sup>7</sup> is H and the other of R<sup>6</sup> and R<sup>7</sup> is halogen, -OR<sup>11</sup>, -NR<sup>11</sup>R<sup>12</sup>, or optionally substituted C<sub>1</sub>-C<sub>6</sub> alkyl.

25. (Original) A method of claim 23, wherein R<sup>8</sup> is halogen.

26. (Original) A method according to claim 1, wherein the carbon atom designated \* is in the R configuration.

27. (Original) A method according to claim 1, wherein the carbon atom designated \* is in the S configuration.

28. (Original) A method comprising a mixture of stereoisomeric compounds of claim 1 wherein the carbon atom designated \* is in the S or R configuration.

29. (Previously presented) A method according to claim 1, wherein the compound is selected from the group consisting of:

2,7-dimethyl-4-phenyl-1,2,3,4-tetrahydroisoquinoline;  
4-(4-methoxy)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
2,7-dimethyl-4-(4-fluoro)phenyl-1,2,3,4-tetrahydroisoquinoline;  
2,7-dimethyl-4-(3-fluoro)phenyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3,4-difluoro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
2,7-dimethyl-4-(4-fluoro-3-methyl)phenyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-chloro-4-fluoro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-chloro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
2,7-dimethyl-4-(4-methyl)phenyl-1,2,3,4-tetrahydroisoquinoline;  
2,7-dimethyl-4-(3-fluoro-4-methyl)phenyl-1,2,3,4-tetrahydroisoquinoline;  
4-(4-chloro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
4-(4-chloro-3-fluoro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3,4-dichloro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
7-ethyl-2-methyl-4-phenyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3,4-difluoro)phenyl-7-ethyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
7-fluoro-4-(4-methoxy)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
7-fluoro-4-(3-fluoro-4-methoxy)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
7-fluoro-4-(3-fluoro-4-methyl)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
7-fluoro-4-(4-chloro-3-fluoro)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3,4-difluoro)phenyl-7-fluoro-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-chloro)phenyl-7-fluoro-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
7-cyano-2-methyl-4-phenyl-1,2,3,4-tetrahydroisoquinoline;  
2-methyl-4-phenyl-7-trifluoromethyl-1,2,3,4-tetrahydroisoquinoline;  
4-phenyl-1,2,7-trimethyl-1,2,3,4-tetrahydroisoquinoline;  
4-(4-chloro)phenyl-1,2-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3,4-difluoro)phenyl-1,2-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
4-phenyl-2,7,8-trifluoromethyl-1,2,3,4-tetrahydroisoquinoline;  
2,7-dimethyl-8-fluoro-4-phenyl-1,2,3,4-tetrahydroisoquinoline;  
2,8-dimethyl-7-fluoro-4-phenyl-1,2,3,4-tetrahydroisoquinoline;  
2,7-dimethyl-8-methoxy-4-phenyl-1,2,3,4-tetrahydroisoquinoline;  
2,7-dimethyl-8-hydroxy-4-phenyl-1,2,3,4-tetrahydroisoquinoline;

2-methyl-4-phenyl-7-trifluoromethoxy-1,2,3,4-tetrahydroisoquinoline;  
4-(3,4-difluoro)phenyl-7-methoxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(4-fluoro-3-methyl)phenyl-7-methoxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-fluoro-4-methyl)phenyl-7-methoxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
7-methoxy-4-(3-methyl)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
2-methyl-7-phenoxy-4-phenyl-1,2,3,4-tetrahydroisoquinoline;  
7-(4-methoxy)phenoxy-2-methyl-4-phenyl-1,2,3,4-tetrahydroisoquinoline;  
7-benzyloxy-2-methyl-4-phenyl-1,2,3,4-tetrahydroisoquinoline;  
7-hydroxy-2-methyl-4-(3-methyl)phenyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-fluoro-4-methyl)phenyl-7-hydroxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(4-fluoro-3-methyl)phenyl-7-hydroxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3,4-difluoro)phenyl-7-hydroxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-cyano)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
2,8-dimethyl-4-phenyl-1,2,3,4-tetrahydroisoquinoline;  
2,8-dimethyl-4-(4-fluoro)phenyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3,4-difluoro)phenyl-2,8-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3,5-difluoro)phenyl-2,8-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
2,8-dimethyl-4-(3-fluoro)phenyl-1,2,3,4-tetrahydroisoquinoline;  
2,8-dimethyl-4-(4-fluoro-3-methyl)phenyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-chloro-4-fluoro)phenyl-2,8-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3,4-dichloro)phenyl-2,8-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-chloro)phenyl-2,8-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
4-(4-chloro)phenyl-2,8-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
4-(4-chloro-3-fluoro)phenyl-2,8-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
2,8-dimethyl-4-(4-methoxy)phenyl-1,2,3,4-tetrahydroisoquinoline;  
4-(4-cyano)phenyl-2,8-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
2,8-dimethyl-4-(4-trifluoromethyl)phenyl-1,2,3,4-tetrahydroisoquinoline;  
2,8-dimethyl-4-(4-methyl)phenyl-1,2,3,4-tetrahydroisoquinoline;  
2-methyl-8-(N-methylamino)methyl-4-phenyl-1,2,3,4-tetrahydroisoquinoline;  
8-(hydroxy)methyl-2-methyl-4-phenyl-1,2,3,4-tetrahydroisoquinoline;  
2-methyl-4-phenyl-8-sulfonamide-1,2,3,4-tetrahydroisoquinoline;  
2-methyl-8-(N-methyl)sulfonamide-4-phenyl-1,2,3,4-tetrahydroisoquinoline;  
8-methoxy-2-methyl-4-(4-methyl)phenyl-1,2,3,4-tetrahydroisoquinoline;

4-(3,5-difluoro)phenyl-8-methoxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-chloro)phenyl-8-methoxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3,4-dichloro)phenyl-8-methoxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(4-chloro-3-fluoro)phenyl-8-methoxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-chloro-4-fluoro)phenyl-8-methoxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3,5-difluoro)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-chloro-5-fluoro)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3,5-difluoro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-chloro-5-fluoro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;  
2-methyl-4-(3,4,5-trifluoro)phenyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-fluoro)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-fluoro-4-methyl)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(4-fluoro-3-methyl)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3,4-difluoro)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-chloro)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(4-chloro-3-fluoro)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-chloro-4-fluoro)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(3-cyano)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(4-acetanilide)-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
4-(4-chloro)phenyl-4-fluoro-2-methyl-1,2,3,4-tetrahydroisoquinoline;  
(3,5-difluoro)-4-phenyl-1,2,7-trimethyl-1,2,3,4-tetrahydroisoquinoline;  
(8-fluoro-2-methyl-4-phenyl-1,2,3,4-tetrahydro-7-isoquinoliny)-N-methylmethanamine;  
(2-methyl-4-phenyl-7-isoquinoliny)-N-methylmethanamine;  
N-methyl-(2-methyl-4-phenyl-7-isoquinoliny)-N-methylmethanamine;  
8-hydroxy-2-methyl-4-phenyl-1,2,3,4-tetrahydro-7-isoquinolinecarbonitrile;  
(2-methyl-4-phenyl-1,2,3,4-tetrahydro-7-isoquinoliny)methanol;  
2-ethyl-4-phenyl-1,2,3,4-tetrahydroisoquinoline; an oxide thereof, a pharmaceutically acceptable salt thereof, a solvate thereof, or prodrug thereof.

30. (Previously Presented) A method of claim 1, wherein the urinary incontinence is urge, stress, or mixed urinary incontinence.